

AMENDMENTS TO THE CLAIMS

1. (Currently Amended) A method of treating a disorder in an individual, the method comprising administering to an individual ~~in need thereof~~ who has the disorder an effective amount of pirfenidone or a pirfenidone analog; comparing a post-treatment level of stress-activated protein kinase (SAPK) activity in a biological sample from the individual to a pre-treatment level of SAPK activity; and adjusting the dose of the pirfenidone or pirfenidone analog based on the results of the comparison step; wherein the disorder is selected from the group consisting of fibrotic disorder, carcinoma, sarcoma, leukemia, lymphoma, viral infection, inflammatory disorder and TNF-mediated disorder.

2. (Original) The method of claim 1, wherein the post-treatment SAPK activity level is from about 10% to about 40% lower than the pre-treatment SAPK activity level, and wherein the adjusting step comprises administering a second dosage of pirfenidone or pirfenidone analog that is at least about 10% higher than the first dosage of pirfenidone or pirfenidone analog.

3. (Original) The method of claim 1, wherein the biological sample is peripheral blood mononuclear cells.

4. (Cancelled).

5. (Currently Amended) The method of claim [[4]]1, wherein said fibrotic disorder is pulmonary fibrosis, renal fibrosis, liver fibrosis, or heart fibrosis.

6. (Cancelled).

7. (Cancelled).

8. (Cancelled).

9. (Cancelled).

10. (Cancelled).

11. (Cancelled).

12. (Original) The method of claim 1, further comprising administering an effective amount of a Type II interferon receptor agonist.

13. (Original) The method of claim 12, wherein the Type II interferon receptor agonist is IFN- γ .

14. (Original) The method of claim 1, further comprising administering an effective amount of a Type I interferon receptor agonist.

15. (Original) The method of claim 14, wherein the Type I interferon receptor agonist is IFN- α .

16. (Currently Amended) A method of treating a disorder in an individual, the method comprising administering to an individual ~~in need thereof~~ who has the disorder an effective amount of pirfenidone or a pirfenidone analog; comparing a second post-treatment level of stress-activated protein kinase (SAPK) activity in a biological sample from the individual to a first post-treatment level of SAPK activity; and adjusting the dose of the pirfenidone or pirfenidone analog based on the results of the comparison step; wherein the disorder is selected from the group consisting of fibrotic disorder, carcinoma, sarcoma, leukemia, lymphoma, viral infection, inflammatory disorder and TNF-mediated disorder.

17. (Currently Amended) A method of inhibiting a stress-activated protein kinase enzymatic activity in a cell of an individual, the method comprising administering to an individual ~~in need thereof~~ an effective amount of pirfenidone or a pirfenidone analog; comparing a post-treatment level of stress-activated protein kinase (SAPK) activity in a biological sample from the individual to a pre-treatment level of SAPK activity; and adjusting the dose of the pirfenidone or pirfenidone analog based on the results of the comparison step; wherein the individual in need thereof has a disorder selected from the group consisting of fibrotic disorder, carcinoma, sarcoma, leukemia, lymphoma, viral infection, inflammatory disorder and TNF-mediated disorder.

18. (New) The method of claim 1, wherein the carcinoma comprises a solid tumor.

19. (New) The method of claim 1, wherein the sarcoma comprises a solid tumor.

20. (New) The method of claim 1, wherein the leukemia comprises a solid tumor.

21. (New) The method of claim 1, wherein the lymphoma comprises a solid tumor.

22. (New) The method of claim 18, wherein the pirfenidone or pirfenidone analog are administered as adjuvant therapy to a primary carcinoma therapy.

23. (New) The method of claim 19, wherein the pirfenidone or pirfenidone analog are administered as adjuvant therapy to a primary sarcoma therapy.

24. (New) The method of claim 20, wherein the pirfenidone or pirfenidone analog are administered as adjuvant therapy to a primary leukemia therapy.

25. (New) The method of claim 21, wherein the pirfenidone or pirfenidone analog are administered as adjuvant therapy to a primary lymphoma therapy.